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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
09/647,695	04/06/01	FARMER	19374-502

HM22/0906

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EXAMINER

DAVIS, R

ART UNIT	PAPER NUMBER
1651	9

DATE MAILED: 09/06/01

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trad marks

**Office Action Summary**

Application No.

09/647,695

Applicant(s)

FARMER ET AL.

Examiner

Ruth A. Davis

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 16 August 2001.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 1-76 is/are pending in the application.
- 4a) Of the above claim(s) 31-76 is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 1-5 and 7-30 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on \_\_\_\_\_ is: a) ☐ approved b) ☐ disapproved by the Examiner.
- If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

**Priority under 35 U.S.C. §§ 119 and 120**

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- \* See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
- a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

**Attachment(s)**

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO-1449) Paper No(s) 7.
- 4) ☐ Interview Summary (PTO-413) Paper No(s). \_\_\_\_\_.
- 5) ☐ Notice of Informal Patent Application (PTO-152)
- 6) ☐ Other: \_\_\_\_\_.

## DETAILED ACTION

### *Election/Restrictions*

1. Applicant's election without traverse of Group I, claims 1 – 30 in Paper No. 8 is acknowledged. Claims 31 – 76 have been withdrawn from consideration.

### *Claim Rejections - 35 USC § 112*

2. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

3. Claims 12 and 24 – 28 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 12 is drawn to a method for decreasing serum cholesterol however is rendered vague and indefinite for reciting “ $5 \times 10^8$  to  $10^9$ ” because it is not clear if the “ $5 \times$ ” applies to both  $10^8$  and  $10^9$  or just  $10^8$ .

Claim 24 recites the limitation "said fibrin" in line 1. There is insufficient antecedent basis for this limitation in the claim.

Claim 26 recites the limitation "said compost" in line 1. There is insufficient antecedent basis for this limitation in the claim.

Claim 27 and its dependent are drawn to a method for decreasing serum cholesterol however are rendered vague and indefinite because it is not clear if the complexation

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(sequestering or chelating) agent is selected from a metal salt of only calcium or a metal salt of each of the named elements.

***Claim Rejections - 35 USC § 102***

4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

(e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371(c) of this title before the invention thereof by the applicant for patent.

5. Claims 1 – 2, 5, 7, 9, 13, 19 and 29 are rejected under 35 U.S.C. 102(b) as being anticipated by Paul (US 5,531,989).

Applicant claims a method for decreasing serum cholesterol and increasing serum HDL comprising orally administering an effective amount of a composition comprising a viable lactic acid bacteria and a therapeutic composition selected from a cholesterol reducing agent or a bifidogenic oligosaccharide wherein the lactic acid bacteria is selected from non-pathogenic members of Bacillus, Lactobacillus, Sporolactobacillus or Bifidobacterium. Specifically, the Lactobacillus is selected from L. acidophilus, L. casei, L. DDS-1, L. GG, L. rhamnosus, L. plantarum, L. salivarius, or L. sporogenes (B. coagulans); the Bifidobacterium is selected from B. adolescentis, B. animalis, B. bifidum, B. bifidus, B. breve, B. infantis, B. infantus, or B.

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longum; the bifidogenic oligosaccharide is selected from fructo-oligosaccharide, gluco-oligosaccharide or trisaccharide raffinose; the cholesterol reducing agent is one of a statin, a bile sequestering compound, a fiber product capable of binding cholesterol, niacin or aspirin and the composition further comprises a food substance, flavoring, vitamin or mineral.

Paul teaches oral administration of a composition comprising dietary fiber (cholesterol reducing agent), fructooligosaccharide (FOS), and a beneficial human intestinal microorganism (lactic acid bacteria) wherein the microorganism is *Lactobacillus* or *Bifidobacteria* (abstract). Specifically, beneficial microorganisms are disclosed to include *L. acidophilus*, *L. casei*, *L. salivarius*, *L. brevis*, *L. plantarum*, *B. adolescentis*, *B. infantis*, *B. longum* and *B. bifidum* (col.4 line 20-30). The ingredients are combined with juice or water and are taken in dosages of about 20 – 400 mg/kg body weight (col.14 line 5-10). Paul discloses compositions such as this have beneficial effects on cholesterol metabolism resulting in decreased serum cholesterol and increased HDL to LDL ratio (col.2 line 35-40). Specifically, FOS is disclosed to reduce serum cholesterol, improve HDL/LDL ratios and increase bifidobacterium populations (col.6 line 55-68).

Though the reference does not specifically teach administering the composition in a method to decrease serum cholesterol and increase serum HDL, the reference does teach these effects. Furthermore, by administering the composition of Paul, it is inherent that serum cholesterol decreased and serum HDL increased. Moreover, by practicing the method of Paul, one in the art would inherently be decreasing serum cholesterol and increasing serum HDL, as claimed by applicant, because the compositions are the same.

Therefore the reference anticipates the claimed subject matter.

***Claim Rejections - 35 USC § 103***

6. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

7. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

8. Claims 1, 8, 10 – 19 24 – 25 and 30 are rejected under 35 U.S.C. 103(a) as being unpatentable over Paul (US 5,531,989).

Applicant claims a method for decreasing serum cholesterol and increasing serum HDL in a patient comprising administering an effective amount of a composition comprising a viable lactic acid bacteria and a therapeutic composition selected from a cholesterol reducing agent or a bifidogenic oligosaccharide wherein the bifidogenic oligosaccharide is selected from fructo-oligosaccharide, gluco-oligosaccharide or trisaccharide raffinose and the cholesterol reducing agent is one of a statin, a bile sequestering compound, a fiber product capable of binding cholesterol, niacin or aspirin. Specifically, the bifidogenic oligosaccharide is a fructose-

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oligosaccharide comprising polymers of fructose and glucose with a chain length of about 4 – 100 sugar units and the fiber is one of gemfibrozil, fenofibrate, psyllium bran, glucomannan or Jerusalem artichoke flour. The composition contains  $10^5$  –  $10^{10}$  viable bacterium/gram composition, 10 mg – 1 gram of bifidogenic oligosaccharide per gram composition and/or 100 – 500 mg bifidogenic oligosaccharide per gram composition. Administration comprises introducing into the gastrointestinal tract 0.1 – 5 grams/day of the composition,  $10^8$  –  $10^{10}$  viable bacterium per day,  $5 \times 10^8$  –  $10^9$  viable bacterium/day, 10 mg – 20 grams of bifidogenic oligosaccharide per day, 150 mg – 5 grams bifidogenic oligosaccharide per day and 500 mg – 50 grams of fiber per day. Finally, the patient is at risk for atherosclerosis, arterial sclerosis, myocardial infarction, heart attack, diabetes, coronary heart disease, angina pectoris or unstable angina.

Paul teaches oral administration of a composition comprising dietary fiber (cholesterol reducing agent), fructooligosaccharide (FOS), and a beneficial human intestinal microorganism (lactic acid bacteria) wherein the microorganism is Lactobacillus or Bifidobacteria (abstract). Paul discloses compositions such as this have beneficial effects on cholesterol metabolism resulting in decreased serum cholesterol and increased HDL to LDL ratio (col.2 line 35-40). Specifically, FOS is disclosed to reduce serum cholesterol, improve HDL/LDL ratios and increase beneficial bifidobacterium populations (col.6 line 55-68).

Although the reference does not specifically teach administering the composition in a method to decrease serum cholesterol and increase serum HDL, the reference does teach these effects. Furthermore, by administering the composition of Paul, it is inherent that serum cholesterol decreased and serum HDL increased. Moreover, by practicing the method of Paul,

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one in the art would inherently be decreasing serum cholesterol and increasing serum HDL, as claimed by applicant, because the compositions are the same.

Paul does not teach the specific amounts of viable microorganisms of bacterium or bifidogenic oligosaccharide per gram of composition or the administration dosages as claimed by applicant. However, it would have been obvious to one of ordinary skill in the art to optimize volumes of ingredients and effective amounts of bacterium because it was routine practice in the art at the time of the invention. At the time of the invention, one of ordinary skill in the art would have been motivated by Paul to optimize the effective amounts to reduce serum cholesterol and increase serum HDL because of these disclosed known actions of the composition. Moreover, at the time of the invention, one of ordinary skill in the art would have been motivated by Paul and conventional practice to optimize administration dosages and volumes of ingredients with a reasonable expectation of success to reduce serum cholesterol and increase serum HDL.

Paul does not teach the fructooligosaccharide with a chain length of 4 – 100 sugar units. However, at the time of the invention, it would have been obvious to one of ordinary skill in the art to substitute or optimize the size of fructooligosaccharide to enhance a desired effect because it was routine practice in the art at the time of the invention. Moreover, at the time of the invention, one of ordinary skill in the art would have been motivated by conventional practice to optimize the size of the fructooligosaccharide in the method of Paul, with a reasonable expectation of success for decreasing serum cholesterol and increasing serum HDL.

Paul does not teach Jerusalem artichoke flour as the fiber source. However, Paul does teach inulin as the fiber source (abstract). Specifically, Paul teaches Jerusalem artichoke is a rich



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source of inulin (fiber) wherein Bifidobacterium uses it as an energy source (col. 6 line 5-20). At the time of the invention, one of ordinary skill in the art would have been motivated to use Jerusalem artichoke flour because of the fiber (inulin) content contained in Jerusalem artichoke as taught by Paul. One of ordinary skill in the art would have been further motivated to use Jerusalem artichoke flour because it was known to be a source of energy for Bifidobacterium as disclosed by Paul.

Paul does not teach the method wherein the patient is at risk for atherosclerosis, arterial sclerosis, myocardial infarction, heart attack, diabetes, coronary heart disease, angina pectoris or unstable angina. However, because of the disclosed beneficial effects on serum cholesterol and LDL/HDL ratios, it would have been obvious to one of ordinary skill in the art to practice the method of Paul on patients at risk for the aforementioned conditions because high cholesterol was a known symptom/indicator of each of the conditions. At the time of the invention, one of ordinary skill in the art would have been motivated to practice the method of Paul on “at risk patients” with a reasonable expectation for success because of the disclosed benefits of reduced serum cholesterol and improved LDL/HDL ratios.

9. Claims 1 – 4 are rejected under 35 U.S.C. 103(a) as being unpatentable over Paul (US 5,531,989) in view of Fukushima et al. (British Journal of Nutrition, 1995).

Applicant claims a method for decreasing serum cholesterol and increasing serum HDL comprising administering an effective amount of a composition comprising a viable lactic acid bacteria and a therapeutic composition selected from a cholesterol reducing agent or a bifidogenic oligosaccharide wherein lactic acid bacteria is selected from non-pathogenic

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members of *Bacillus*, *Lactobacillus*, *Sporolactobacillus* or *Bifidobacterium*. Specifically, wherein *Bacillus* is selected from *B. coagulans*, *B. coagulans* Hammer, *B. brevis* subspecies *coagulans* or *B. laevolacticus*. More specifically, *B. laevolacticus*.

Paul teaches oral administration of a composition comprising dietary fiber (cholesterol reducing agent), fructooligosaccharide (FOS), and a beneficial human intestinal microorganism (lactic acid bacteria) wherein the microorganism is *Lactobacillus* or *Bifidobacteria* (abstract).

Paul discloses compositions such as this have beneficial effects on cholesterol metabolism resulting in decreased serum cholesterol and increased HDL to LDL ratio (col.2 line 35-40). Specifically, FOS is disclosed to reduce serum cholesterol, improve HDL/LDL ratios and increase *bifidobacterium* populations (col.6 line 55-68).

Although the reference does not specifically teach administering the composition in a method to decrease serum cholesterol and increase serum HDL, the reference does teach these effects. Furthermore, by administering the composition of Paul, it is inherent that serum cholesterol decreased and serum HDL increased. Moreover, by practicing the method of Paul, one in the art would inherently be decreasing serum cholesterol and increasing serum HDL, as claimed by applicant, because the compositions are the same.

Paul does not teach the method wherein *Bacillus* species are used. However, at the time of the invention, it would have been obvious to one of ordinary skill in the art to utilize a *Bacillus* in the methods of Paul because Fukushima et al. (1995) teach the effects of a probiotic composed of *Bacillus* include reduced serum cholesterol and increased serum HDL (abstract). Though Fukushima et al. does not teach specific species of *Bacillus*, the reference does teach the effect is achieved with the genus. Therefore, at the time of the invention, it would have been

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obvious to one of ordinary skill in the art to utilize any *Bacillus* as they were known to have the claimed effect. Moreover, at the time of the invention, one of ordinary skill in the art would have been motivated by Fukushima to use a *Bacillus* in the methods of Paul with a reasonable expectation for success of decreasing serum cholesterol and increasing serum HDL because of the disclosed properties of *Bacillus* specifically achieving the claimed effect.

10. Claims 1 and 19 – 25 are rejected under 35 U.S.C. 103(a) as being unpatentable over Paul (US 5,531,989) in view of Bova et al. (US Application 2001/0006644 A1).

Applicant claims a method for decreasing serum cholesterol and increasing serum HDL comprising administering an effective amount of a composition comprising a viable lactic acid bacteria and a therapeutic composition selected from a cholesterol reducing agent or a bifidogenic oligosaccharide wherein the cholesterol reducing agent is one of a statin, a bile sequestering compound, a fiber product capable of binding cholesterol, niacin or aspirin. Specifically, the statin is one of cervastatin, fluvastatin, lovastatin, pravastatin or simvastatin with administration of 10 – 80 mg of statin per day, the bile sequestering agent is one of colestipol or cholestyramine with administration of 1 – 20 grams of bile sequestering agent per day and the fiber is one of gemfibrozil, fenofibrate, psyllium bran, glucomannan or Jerusalem artichoke flour with administration of 500 mg – 50 grams of fiber per day.

Paul teaches oral administration of a composition comprising dietary fiber (cholesterol reducing agent), fructooligosaccharide (FOS), and a beneficial human intestinal microorganism (lactic acid bacteria) wherein the microorganism is *Lactobacillus* or *Bifidobacteria* (abstract). Paul discloses compositions such as this have beneficial effects on cholesterol metabolism

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resulting in decreased serum cholesterol and increased HDL to LD ratio (col.2 line 35-40).

Specifically, FOS is disclosed to reduce serum cholesterol, improve HDL/LDL ratios and increase bifidobacterium populations (col.6 line 55-68).

Though the reference does not specifically teach administering the composition in a method to decrease serum cholesterol and increase serum HDL, the reference does teach these effects. Furthermore, by administering the composition of Paul, it is inherent that serum cholesterol decreased and serum HDL increased. Moreover, by practicing the method of Paul, one in the art would inherently be decreasing serum cholesterol and increasing serum HDL, as claimed by applicant, because the compositions are the same.

Paul does not teach the method wherein therapeutic agents include statins, cervastatin, flucastatin, lovastatin, pravastatin or simvastatin; bile sequestering agents, colestipol or cholestyramine; fiber sources gemfibrozil or fenofibrate; niacin or aspirin wherein they are administered in amounts as claimed by applicant. However, at the time of the invention, one of ordinary skill in the art would have been motivated to include any of the aforementioned agents in the composition and method of Paul because Bova et al. teach each of these are known to lower serum cholesterol (p.1, paragraphs 0006 – p.2 paragraph 0009). Bova et al. teach numerous methods for reducing serum cholesterol and increasing HDL cholesterol levels have been proposed to include administration of hypolipidemic agents, or lipid altering agents (p.1 paragraph 0006). Specifically, niacin and the named statins are disclosed to reduce total serum cholesterol and increase serum HDL cholesterol (p.2 paragraph 0010, 0017) while bile sequestering agents are disclosed to be a first choice for treating hypercholesterolemia due to their efficacy and proved usefulness (p.3 paragraph 0020). Furthermore, Bova et al. teach the

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administration of statins in dosages between 10 – 80 mg, 5 – 80 mg and 20 – 80 mg (p.4 paragraph 0033). Therefore, at the time of the invention, one of ordinary skill in the art would have been motivated by Bova et al. to include any of the above mentioned agents in the method of Paul, with a reasonable expectation of success for decreasing serum cholesterol and increasing serum HDL because they were well known in the art for reducing cholesterol and increasing HDL as demonstrated by the references cited above.

The above references do not teach administration volumes of bile sequestering agents or fiber. However, at the time of the invention, it would have been obvious to one of ordinary skill in the art to optimize dosages of effective ingredients because it was routine practice in the art at the time of the invention. Moreover, at the time of the invention, one of ordinary skill in the art would have been motivated by conventional practice to optimize dosages of effective ingredients with a reasonable expectation of success for decreasing serum cholesterol and increasing serum HDL.

11. Claims 1 and 26 – 28 are rejected under 35 U.S.C. 103(a) as being unpatentable over Paul (US 5,531,989) in view of Mandeville, III et al. (US 5,607,669).

Applicant claims a method for decreasing serum cholesterol and increasing serum HDL comprising administering an effective amount of a composition comprising a viable lactic acid bacteria and a therapeutic composition selected from a cholesterol reducing agent or a bifidogenic oligosaccharide wherein the composition further comprises a cholic acid complexation agent selected from a metal salt of calcium, chromium copper, iodine, iron,

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magnesium, manganese, potassium, sodium or zinc. Specifically, the metal salt is calcium citrate, potassium gluconate, magnesium citrate or chromium picollinate.

Paul teaches oral administration of a composition comprising dietary fiber (cholesterol reducing agent), fructooligosaccharide (FOS), and a beneficial human intestinal microorganism (lactic acid bacteria) wherein the microorganism is Lactobacillus or Bifidobacteria (abstract). Paul discloses compositions such as this have beneficial effects on cholesterol metabolism resulting in decreased serum cholesterol and increased HDL to LD ratio (col.2 line 35-40). Specifically, FOS is disclosed to reduce serum cholesterol, improve HDL/LDL ratios and increase bifidobacterium populations (col.6 line 55-68).

Although the reference does not specifically teach administering the composition in a method to decrease serum cholesterol and increase serum HDL, the reference does teach these effects. Furthermore, by administering the composition of Paul, it is inherent that serum cholesterol decreased and serum HDL increased. Moreover, by practicing the method of Paul, one in the art would inherently be decreasing serum cholesterol and increasing serum HDL, as claimed by applicant, because the compositions are the same.

Paul does not teach the method further comprising administering a cholic acid complexation agent. However, at the time of the invention, one of ordinary skill in the art would have been motivated to include a cholic acid complexation (or sequestering, chelating) agent because Mandeville, III et al. teach that it is advantageous to sequester primary bile acids to significantly increase the reduction of serum lipid levels (col.1 line 55-65). At the time of the invention, it was known in the art that cholic acid is a primary bile acid and that sorption of bile acids is related to decreasing serum cholesterol (US 5427777 col.2 line 25-35). Though the

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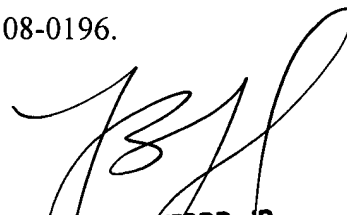
reference does not teach the specific cholic acid complexation (sequestering) agents claimed by applicant, it would have been obvious to one of ordinary skill in the art to use any complexing agent as they were known in the art at the time of the invention. Moreover, at the time of the invention, one of ordinary skill in the art would have been motivated by Mandeville, III et al. to include a cholic acid complexation (sequestering) agent in the method of Paul with a reasonable expectation of success for reducing serum cholesterol because of the cholesterol/lipid reducing effects as demonstrated by Mandeville, III et al.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ruth A. Davis whose telephone number is 703-308-6310. The examiner can normally be reached on M-H (7:00-4:30); altn. F (7:00-3:30).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Wityshyn can be reached on 703-308-4743. The fax phone numbers for the organization where this application or proceeding is assigned are 703-308-4242 for regular communications and 703-308-4242 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0196.

Ruth A. Davis  
September 5, 2001



LEON B. LANKFORD, JR.  
PRIMARY EXAMINER